REIMBURSABLE DETAIL OPPORTUNITY

CENTER FOR TOBACCO PRODUCTS

The Center for Tobacco Products is offering a Detail Opportunity for a position as **Senior Regulatory Counsel, GS-0301-15**. The Detail is available immediately for a period up to 90 days. Commissioned Corps officers are encouraged to apply.

No. of Positions:	One
Bargaining Unit Status:	Non-Bargaining Unit Position
Ethics Requirements:	Ethics pre-clearance is required for this position
Office Duty Location:	FDA Center for Tobacco Products Office of Compliance and Enforcement 10903 New Hampshire Ave. Silver Spring, MD 20993
Opening Date:	**March 19, 2018**
Closing Date:	**March 23, 2018**
Area of Consideration:	Open to all career/career-conditional FDA-employees

On June 22, 2009, the President signed into law the Family Smoking Prevention and Tobacco Control Act (the Tobacco Control Act) (Public Law 111-31). The Tobacco Control Act granted FDA important new authority to regulate the manufacturing, marketing, and distribution of tobacco products to protect the public health generally and to reduce tobacco use by minors. To carry out this responsibility, FDA established the Center for Tobacco Products (CTP).

Major Duties:

The employee performs duties that include resolving a broad range of issues such as:

- Plan long term project related to Tobacco Product Manufacturing Practice regulation, including developing timelines and schedules for review, internal and external briefing, TPSAC, and public hearings
- Oversee efforts to advance rulemaking process through publication and public comment period
- Utilize expertise in manufacturing practice toward development and enhancement of proposed regulation

Skills:

- Knowledge of enabling tobacco legislation, policies, implementing regulations and procedures, policies and guidelines, and interrelationships of compliance organizations and programs
- Ability to recognize the need for and develop new procedures to solve critical or novel problems or perform more refined analyses
- Skill in collaborating with experts in a variety of disciplines on legal, and regulatory issues, as well as excellent interpersonal skills
- Experience writing a variety of regulatory and policy documents that require conducting research on regulatory issues and interpreting issues regarding regulations and policies that affect the operations of a regulatory program

Application Process:

Supervisory concurrence is required to accept a detail; it is **NOT** required to apply.

This Detail opportunity is open to:

- Qualified candidates at the GS-15 grade level
- Public Health Service Commissioned Corps Officers.

Interested applicants should submit via email a resume, SF-50 and statement of interest to:

Anne Gentilcore and Michele Quander
Office of Management
Center for Tobacco Products, FDA
anne.gentilcore@fda.hhs.gov | Michele.quander@fda.hhs.gov

Question/s about the position, please contact Jesse Hardin at 301-796-6830.

Travel Expenses will not be paid.

Applications/resumes must be submitted by March 23, 2018.

This is not an official vacancy announcement under the Merit Promotion System.